Attorney's Docket No. 007157/270549

ENTE TRADEN

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Snyder et al.

Confirmation No.:

4831

Appl. No.:

10/690,462

Group Art Unit:

1614

Filed:

October 21, 2003

Examiner:

(not yet assigned)

For:

CURCUMIN ANALOGS WITH ANTI-TUMOR AND

ANTI-ANGIOGENIC PROPERTIES

Mail Stop Missing Parts Commissioner for Patents

May 17, 2004

P.O. Box 1450

Alexandria, VA 22313-1450

INFORMATION DISCLOSURE STATEMENT CITATION UNDER 37 C.F.R. § 1.97

Attached is a list of documents on Form PTO-1449. It is requested that the Examiner consider these documents and officially make them of record in accordance with the provisions of 37 C.F.R. § 1.97 and Section 609 of the MPEP. By submitting the listed documents, Applicant in no way makes any admission as to the prior art status of the listed documents, but is instead submitting the listed documents for the sake of full disclosure.

All documents were supplied in, or cited by the Office during prosecution of, parent Application No. 09/729,662 filed December 4, 2000. Since the benefit of this application was claimed under 35 U.S.C. 120, no copies need to be furnished in accordance with 37 C.F.R. 1.98(d); however, copies will be furnished on request.

Respectfully submitted,

Registration No. 47,683

Customer No. 00826 ALSTON & BIRD LLP

Bank of America Plaza 101 South Tryon Street, Suite 4000 Charlotte, NC 28280-4000

Tel Raleigh Office (919) 862-2200

Fax Raleigh Office (919) 862-2260

"Express Mail" mailing label number: EV387076958US

Date of Deposit: May 17, 2004

I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above and is addressed to: Mail Stop Missing Parts, Commissioner for Patents, P.O., Box 1450, Alexandria, VA 22313-1450

Tracey S. Wrig

OIPE

MAY 1 7 2004

Substitute for form	1449/PTO
(Revised 04/2003)	

INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Sheet	1 1	of	3

Complete if Known				
Application Number	10/690,462			
Filing Date	October 21, 2003			
First Named Inventor	Snyder			
Group Art Unit	1614			
Examiner Name	(not yet assigned)			
Attorney Docket Number	007157/270549			

	U. S. PATENT DOCUMENTS						
Examiner Initials*	Cite No.	Document Number Number - Kind Code (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages of Relevant Figures Appear		
	1	US-3,114,775	12-17-1963	Hughes et al.			
	2	US-3,911,129	10-07-1975	Krapcho et al.			
	3	US-4,127,667	11-28-1978	Rovnyak			
	4	US-4,415,621	11-15-1983	Specht et al.			
	5	US-4,755,450	07-05-1988	Sanders et al.			
	6	US-4,987,057	01-22-1991	Kaji et al.			
	7	US-5,700,804	12-23-1997	Collins et al.			
	8	US-5,811,218	09-22-1998	Kaji et al.			
	9	US-5,852,018	12-22-1998	Bryans et al.			
	10	US-6,022,597	02-08-2000	Yan et al.			
•	11	US-2002/0006966 A1	01-17-2002	Arbiser			

		FORI	EIGN PATENT I	DOCUMENTS		
Examiner Initials	Cite No.	Foreign Patent Document Country Code - Number Kind Code (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	English Language Translation Attached
	12	JP - 03/44643	02-26-1991	Hioki et al.		
	13	WO – 01/46110	06-28-2001	The University of Georgia Research Foundation, Inc. et al.		
			·			

Examiner	Date	
Signature	Considered	

^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

RTA01/2144500v1

Substitute for form 1449/PTO (Revised 04/2003)

INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use ar many sneets as necessary)

Complete if Known				
Application Number	10/690,462			
Filing Date	October 21, 2003			
First Named Inventor	Snyder			
Group Art Unit	1614			
Examiner Name	(not yet assigned)			
Attorney Docket Number	007157/270549			

MAY 1 7 2004 00

/	£	OTHER DOCUMENTS	
Examiner Initials	Cite No.	magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	English Language Translatio Attached
_	14	ARTICO, et al., "Geometrically and Conformationally Restrained Cinnamoyl Compounds as Inhibitors of HIV-1 Integrase: Synthesis, Biological Evaluation, and Molecular Modeling," J. Med. Chem., 1998, pp. 3948-3960, Vol. 41, No. 21.	
	15	CREMLYN et al., "The Synthesis and Chlorosulfonation of Some Diarylidene and Heteroarylidene Ketones with Varying Alicyclic Ring Size", Phosphorus, Sulfur, and Silicon, 1995, pp. 205-217, Vol. 107.	
	16	DINKOVA-KOSTOVA, et al., "Chemoprotective Properties of Phenylpropenoids, Bis(benzylidene)cycloalkanones, and Related Michael Reaction Acceptors: Correlation of Potencies as Phase 2 Enzyme Inducers and Radical Scavengers," J. Med. Chem., 1998, pp.5287-5296, Vol. 41, No. 26.	
	17	EL-SUBBAGH, et al., "Synthesis and Biological Evaluation of Certain α,β-Unsaturated Ketones and Their Corresponding Fused Pyridines as Antiviral and Cytotoxic Agents," J. Med. Chem., 2000, pp.2915-2921, Vol. 43, No. 15.	
-	18	FUJISAKI, et al., JP 62225562, 1988 (CA 108:77360).	
	19	GUTKOWSKA, et al., Acta Poloniae Pharmaceutica, 1985, pp. 437-441, Vol. 42, No. 5 (CA 107:115819).	
	20	GUTKOWSKA, et al., Acta Poloniae Pharmaceutica, 1989, pp. 212-218, Vol. 46, No. 3 (CA 112:216649).	
	21	HAMMAM, et al., "Synthesis and Anti-Cancer Activity of Pyridine and Thiazolopyrimidine Derivatives Using 1-Ethylpiperidone as a Synthon," Indian J. Chem., 2001, pp. 213-221, Vol. 40B.	
	22	KEINAN, et al., J. Org. Chem., 1983, pp. 5302-5309, Vol. 48, No. 26.	
	23	LI, et al., "Samarium (III) Iodide Promoted Preparation of α,α' – bis(substituted benzylidene) cyclohexanones from Benzaldehydes and Cyclohexanone," J. Chem, Research (S), 2000, pp. 580-581.	
	24	MAHFOUZ, et al., "Synthese mehrfach oxigenierter 2-Hydroxyxanthone," Arch. Pharm. (Weinheim), 1990, pp. 163-169, Vol. 323.	
	25	NAKANO, et al., "A Convenient Synthesis of α,α' – Bis(substitutedbenzylidene)cycloalkanones," Chemistry Letters, 1993, pp. 2157-2158.	

Examiner	Date	
Signature	Considered	

^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

RTA01/2144500v1

		6 LPE				
	r form 1449/	го	SC.		Complete if Known	
(Revised 04/	(2003)	MAY 1 7 200	14 a	Application Number	10/690,462	
INDOD				Filing Date	October 21, 2003	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as RADE)			SORE	First Named Inventor	Snyder	
STATE	MENT B	PADE	MAN'T	Group Art Unit	1614	
(U.	se as many she	ets as its estais)		Examiner Name	(not yet assigned)	
Sheet	3	of	3	Attorney Docket Number	007157/270549	

	OTHER DOCUMENTS					
Examiner Initials	Cite No.					
	26	OJIMA, et al., Bull. Chem. Soc. Jpn., 1977, pp. 1522-1526, Vol. 50, No. 6 (CA 87:20055).				
	27	PIVNENKO, et al., Zh. Org. Khim, 1972, pp. 1096-1102, Vol. 42, No. 5 (CA 84:513251).				
	28	PIVNENKO, et al., Zh. Org. Khim, 1975, pp. 2527-2533, Vol. 11, No. 12 (CA 84:73234).				
	29	SHOPPEE, et al., J. Chem. Soc. Perkin Trans I, 1977, pp. 1028-1030, Vol. 9 (CA 87:102029).				
<i>✓</i>	30	SUN, et al., "Design, Synthesis, and Evaluations of Substituted 3-[(3- or 4-Carboxyethylpyrrol-2-yl)methylidenyl]indolin-2-ones as Inhibitors of VEGF, FGF, and PDGF Receptor Tyrosine Kinases," J. Med. Chem., 1999, pp. 5120-5130, Vol. 42, No. 25.				
	31	SUN, et al., "Identification of Substituted 3-[(4,5,6,7-Tetrahydro-1 <i>H</i> -indol-2-yl)methylene]-1,3-dihydroindol-2-ones as Growth Factor Receptor Inhibitors for VEGF-R2 (Flk-1/KDR), FGF-R1, and PDGF-Rβ Tyrosine Kinases," <i>J. Med. Chem.</i> , 2000, pp. 2655-2663, Vol. 43, No. 14.				
	32	SUN, et al., "Synthesis and Biological Evaluations of 3-Substituted Indolin-2-ones: A Novel Class of Tyrosine Kinase Inhibitors That Exhibit Selectivity toward Particular Receptor Tyrosine Kinases," J. Med. Chem., 1998, pp. 2588-2603, Vol. 41, No. 14.				
	33	TEUSCHER, "Potentiell antiangiogene Substanzen aus der Gruppe der α, α'-Bis(amidinobenzyl)cycloalkanon-Derivate und α-(Arylsulfonylamino)-ω-phenylcarbonsäure-4-amidinoanilide," <i>Pharmazie</i> , 1987, pp. 109-110, Vol.42, H.2.				
	34	THALOOR, et al., "Inhibition of Angiogenic Differentiation of Human Umbilical Vein Endothelial Cells by Curcumin," Cell Growth & Differentiation, 1998, pp. 305-312, Vol. 9.				
	35	VIETH, et al., "DoMCoSAR: A Novel Approach for Establishing the Docking Mode That Is Consistent with the Structure-Activity Relationship. Application to HIV-1 Protease Inhibitors and VEGF Receptor Tyrosine Kinase Inhibitors", J. Med. Chem., 2000, pp. 3020-3032, Vol. 43, No. 16.				
	36	WIEMER et al., "Vidalols A and B, New Anti-Inflammatory Bromophenols from the Caribbean Marine Red Alga Vidalia obtusaloba," Experientia, 1991, pp. 851-853, Vol. 47.				
	37	ZHENG, et al., Zhongguo Yiyao Gonye Zazhi, 1997, p. 230231, Vol. 28, No. 5 (CA 115:102878).				

Examiner	Date	
Signature	Considered	

^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

RTA01/2144500v1